Cancer and
Advance Care Planning

You’ve been diagnosed with cancer. Now what?
You have a lot to think about and it can be difficult to know where to start. One important thing you should think about is your wishes for your care in case you can’t speak for yourself. This is called advance care planning.

**Benefits of Advance Care Planning:**

- Peace of mind for your loved ones
- Knowing your voice is heard
- A gift to your family

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Advance care planning is:

• An opportunity for you to reflect on your values, beliefs and wishes for care;

• Conversations with family and friends to let them know your future health and personal care wishes in case you become unable to speak for yourself;

• A way to give your loved ones peace of mind in making decisions about your care and treatment during a difficult time.

You may never need an advance care plan but if you do, your family and other caregivers will appreciate knowing what you would want done next.
You might have a number of questions during the course of your cancer treatment. It’s important to discuss them with your doctor and other members of your cancer care team so that you can make informed decisions about your Advance Care Plan:

- What are your options if your health quickly gets worse?
  - How will the treatment plan start? Is the treatment meant to cure the cancer or just make you more comfortable or both?
  - What side effects or complications could be caused by your treatment?

Advance care planning is like retirement planning — it is important to start early, even if you don’t need it for many years to come.

– Health Care Provider

You might have a number of questions during the course of your cancer treatment. It’s important to discuss them with your doctor and other members of your cancer care team so that you can make informed decisions about your Advance Care Plan:

- What impact will cancer have on your life?
  - What is the likely course?

What are your options if your health quickly gets worse?

- How will the treatment plan start? Is the treatment meant to cure the cancer or just make you more comfortable or both?
- What side effects or complications could be caused by your treatment?
What is the best you can hope for with this treatment?

- Are you eligible for a **clinical trial** or experimental treatment?
- How will you know if the cancer has progressed? Will there be any signs or **symptoms**?
- How do we make decisions about whether you should be in hospital or at home?
- Does having advanced cancer change the effectiveness of any other **treatments** you might have in an intensive care unit?

What options are available to you if you don’t want to start or continue treatment?

- What are your options if your health quickly gets worse?
- When would someone consider palliative care? What palliative care services are available in your community?
- What about **life prolonging** or **life sustaining treatments**? How do you decide whether or not to have these **treatments**?

Make sure that you ask these questions throughout your cancer treatment, not just at the beginning. It’s important to tell your health care team how much information you would like and what is important to you.
Advance Care Planning

When you are capable of speaking for yourself, health care providers will always speak directly with you. But, should the time come when you cannot speak for yourself, your substitute decision maker and/or your advance care plan will make sure that your voice is heard.

How Do You Plan?

• **Think** about what’s right for you: What are your values, wishes, beliefs and understanding about your care and specific medical treatments?

• **Learn** about medical procedures: Some may improve your quality of life; others may only keep you alive longer.

• **Decide** on a substitute decision-maker: Choose someone who would honour and follow your wishes if you can’t speak for yourself.

• **Talk** about your wishes: It is important to discuss your wishes with your loved ones, your family physician and your cancer care team.

• **Record** your wishes: It’s a good idea to write down or make a recording of your wishes.

Talking with my family wasn’t easy for me. I feel more at peace now that they all know what I want if the day comes when I can’t tell them.

– Patient
The following questions can help you identify personal goals, wishes and values to guide your discussions and decisions:

What is most important to you about your health and the care you receive right now?

- What makes each day worthwhile?
- What makes you happy?
- How do your decisions about your cancer, your care and your treatment affect your loved ones? Does this change the way you feel about treatment?
- Do you have fears or worries regarding your treatment?
- Do you have a preference regarding location of care (for example, at home, in a health care institution) if your cancer gets worse?
- Is there anything that feels “undone” about your life?
- Who can you rely on to help you through any challenges?
- Do your religious, cultural or personal beliefs affect your decisions?
I wish we had made a plan earlier. Things happened really fast, and by the time he was on a ventilator, it was difficult to know what he would have wanted.

– Family member
Where can you go for more information about cancer and Advance Care Planning?

**Advance Care Planning**
www.advancecareplanning.ca
This website has lots of information and tools to help you make a plan.

The Speak Up Workbook can help you explore your values and write down your wishes for care so that you can share them with others. You can also download a wallet card so that others will know who your substitute decision-maker is.

**American Society of Clinical Oncology**
Advanced Cancer Care Planning: What Patients and Families Need to Know About Their Choices When Facing Serious Illness
http://www.cancer.net/coping/advanced-cancer-care-planning

**Cancer Help UK**
Decisions About Your Care at the End of Life

**National Cancer Institute**
Coping with Advanced Cancer

Members of your cancer care team, including counsellors, social workers, nurses and doctors, can also tell you about resources in your community, province or territory.

What could happen if you don’t make an advance care plan?

- Your loved ones may not be sure what you would want.
- Your loved ones might disagree on what care you would want.
Definition of Terms

**Advanced cancer:** Cancer that cannot be cured with treatment.

**Advance Care Plan (ACP):** A verbal or written plan to let others know what kind of health and personal care you want if you become incapable of consenting to or refusing treatment or other care. An ACP is based on a process of reflection and considers your values, wishes and beliefs.

**Clinical trial:** The study of new cancer treatments.

**Cure:** The treatment causes the cancer to be eliminated from the body.

**Health care provider/health care professional:** A person licensed, certified or registered in their province or territory to provide health care, such as doctors, nurses and social workers.

**Life sustaining/life prolonging treatment:** A medical treatment that is not meant to cure the cancer but to support how the body is working, such as CPR, fluids, breathing machines or antibiotics.
**Oncologist**: A doctor who specializes in treating cancer. Some oncologists specialize in a particular type of cancer treatment. For example, a radiation oncologist specializes in treating cancer with radiation.

**Palliative Care**: Palliative care focuses on providing good quality of life – in other words, keeping you as comfortable and free of pain and other distress as possible. Palliative care may involve medicines, treatments, physical care, psychological/social services and spiritual support for you, your family and others who are helping to care for you.

**Progression (of cancer)**: Cancer spreads or grows larger in the body.

**Recurrence**: Cancer that has returned, usually after a period of time during which the cancer could not be detected. The cancer may come back to the same body part as the original (primary) tumour, or it can affect another part of the body. Recurrence can also be called “recurrent cancer”.

**Remission**: A decrease in or disappearance of signs and symptoms of cancer. In partial remission, some, but not all, signs and symptoms of cancer have disappeared. In complete remission, all signs and symptoms of cancer have disappeared, although cancer may still be in the body.

**Side effects**: Physical or psychological effects of the treatment or the cancer itself. For example, side effects can include pain, tiredness, nausea, depression or anxiety.

**Symptoms**: Signs that you are unwell. For example, pain, vomiting, loss of appetite or high fever can be signs that something is wrong.

**Tumour**: An abnormal mass of tissue in the body that happens when cells divide more than they should or do not die when they should. Tumours may be benign (not cancer) or malignant (cancer), and they can also be called a neoplasm.

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1 Definitions adapted from Speak Up (www.advancecareplanning.ca) and National Cancer Institute (www.cancer.gov/dictionary)
This tool was developed in collaboration with the Advance Care Planning in Canada initiative found at www.advancecareplanning.ca, the BC Cancer Agency and members of the BC Patient Voices Network.

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The BC Cancer Agency, an agency of the Provincial Health Services Authority, provides a province-wide, population-based cancer control program for the residents of British Columbia and the Yukon. The Agency accepts patients who have been diagnosed with cancer and are referred by a physician.

Advance Care Planning in Canada and the Speak Up campaign are managed by the Canadian Hospice Palliative Care Association (CHPCA), a national, bilingual charitable non-profit association with membership comprised of individuals and hospice palliative care programs and services from every province and territory. For more information and advance care planning resources, please visit: www.advancecareplanning.ca